## Wyoming Department of Health Institutional Review Board

## RESEARCH THAT DOES NOT INVOLVE DIRECT HUMAN SUBJECT PARTICIPATION APPLICATION

Attach this application to a WDH-IRB Table of Elements for Protocol Summary and submit to the IRB for approval prior to collection or release of any data or material.

SPECIFICS OF USE OF EXISTING DATA OR BIOLOGICAL MATERIALS	
1.	State why & how the existing data or biological materials were collected, or how data or biological materials to be used in this research will be collected for non-research purposes.
2.	Were the existing data or specimens originally stored in a way that could reveal the identity of the person from whom the material originated? YES NO
3.	Will the research records carry any identifiers that could link the information to the person from whom the material originated? YES NO (If yes discontinue application, submit protocol for IRB Review)
4.	What type of data and/or biological specimens will be used for research?
5.	From how many persons did/will the data or biological specimens originate?
6.	From what source(s) will the data or biological specimens be procured?
7.	How will the investigators gain access to the data or biological specimens?

8. If the data or biological specimens were originally collected for non-research purposes, have the persons from whom the material originated agreed that the material might also be used

for research purposes? YES NO

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9.	Does use of the data or biological specimens involve information, which if revealed, could place someone at risk of criminal or civil liability, or be damaging to their financial standing, employability or reputation? YES NO
10	. How will the permission of the persons, from whom the data or biological specimens originated, be obtained to use the material for research purposes?
11.	. What measures will be taken to keep the research records confidential?
12.	Describe any access that researchers will have to information that is not essential to the research, what will be done with this non-essential information, and how it will be protected.
13.	. Could the research to be conducted on the data or biological specimens reveal information of potential benefit to the persons from whom the material originated? YES NO (If YES, describe plans to inform participants about their rights)
14	Could the research lead to the development of a commercial product that may bring financial benefit to the investigators and/or the sponsor? YES NO (If YES, describe plans to inform participants about their rights)
IR	B#: (assigned by the WDH IRB)
Stı	udy Title
Da	te of submission of this form
Pr	incipal Investigator

**Principal Investigator Signature**